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(FILE 'HOME' ENTERED AT 08:23:52 ON 27 NOV 2007)

FILE 'CAPLUS' ENTERED AT 08:24:07 ON 27 NOV 2007 E US2006-566485/APPS

L1 1 S E3

FILE 'REGISTRY' ENTERED AT 08:25:13 ON 27 NOV 2007

FILE 'CAPLUS' ENTERED AT 08:25:26 ON 27 NOV 2007
L2 TRA L1 1- RN : 3 TERMS

FILE 'REGISTRY' ENTERED AT 08:25:26 ON 27 NOV 2007

L3 3 SEA L2

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L4 1 S E3

FILE 'REGISTRY' ENTERED AT 08:30:01 ON 27 NOV 2007

STR 182878-97-3

L6 1 S L5 EXA FUL

FILE 'CAPLUS' ENTERED AT 08:30:30 ON 27 NOV 2007

L7 3 S L6

L8 1 S L7 AND EMOLLIENT

L9 0 S L7 AND PETROLATUM

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L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:161002 CAPLUS

DOCUMENT NUMBER: 142:246185

TITLE: Pharmaceutical compositions containing quinazoline

derivatives

INVENTOR(S):
Kriwet, Katrin

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

SOURCE: PCT Int. Appl., 15 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE: Englis
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.							APPLICATION NO.										
								WO 2004-EP9273										
WO 2005016322												•						
	W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑŻ,	BA,	BB,	ВG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
	¥.	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	ΚP,	KR,	KZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NA,	NI,	
		NO,	NΖ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
		ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	
		ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	
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		SI,	SK,	TR,	BF,	ΒJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	
			TD,															
AU 2004264680															0040			
CA 2534492			A1		2005	0224		CA 2	004-	2534	492			0040				
									AT 2004-764259									
EP	1675	621			A2		20060705 EP 2004-764259					59	20040818					
EP	1675																	
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OTHER S	OURCE	(S):			MAR	PAT	142:	2461	85					-				

10/566485

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YOU HAVE REQUESTED DATA FROM 3 ANSWERS - CONTINUE? Y/(N):y

L3 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2007 ACS on STN

RN 182878-97-3 REGISTRY

ED Entered STN: 07 Nov 1996

CN Quinazoline, 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl- (CA INDEX NAME) OTHER NAMES:

CN 6-[2-(2,5-Dimethoxyphenyl)ethyl]-4-ethylquinazoline

MF C20 H22 N2 O2

SR CA

LC STN Files: CA, CAPLUS, CASREACT, IMSDRUGNEWS, IMSRESEARCH, PROUSDDR, SYNTHLINE, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 3 REFERENCES IN FILE CA (1907 TO DATE)
- 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d l7 ibib abs hitstr

ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:161002 CAPLUS

DOCUMENT NUMBER: 142:246185

TITLE: Pharmaceutical compositions containing quinazoline

derivatives

INVENTOR (S): Kriwet, Katrin

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

PCT Int. Appl., 15 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

F	PATENT NO.							APPLICATION NO.					DATE						
WO 2005016322 WO 2005016322				A2 20050224			0224	WO 2004-EP9273						20040818					
***		W:	AE, CN, GE, LK, NO, TJ, BW, AZ, EE,	AG, CO, GH, LR, NZ, TM, GH, BY, ES,	AL, CR, GM, LS, OM, TN, GM, KG, FI,	AM, CU, HR, LT, PG, TR, KE, KZ, FR,	AT, CZ, HU, LU, PH, TT, LS, MD, GB,	AU, DE, ID, LV, PL, TZ, MW, RU, GR,	AZ, DK, IL, MA, PT, UA, MZ, TJ,	DM, IN, MD, RO, UG, NA, TM, IE,	DZ, IS, MG, RU, US, SD, AT,	BG, EC, JP, MK, SC, UZ, SL, BE, LU, GA,	EE, KE, MN, SD, VC, SZ, BG, MC,	EG, KG, MW, SE, VN, TZ, CH, NL,	ES, KP, MX, SG, YU, UG, CY, PL,	FI, KR, MZ, SK, ZA, ZM, CZ, PT,	GB, KZ, NA, SL, ZM, ZW, DE, RO,	GD, LC, NI, SY, ZW AM, DK, SE,	
A	U.	20042		TD, 30		A1		2005	0224		AU 2	2004-	2646	80		2	0040	818	
		25344				A1						2004-							
						т		2006	0315		AT 2	2004-	7642	59		2	0040	818	
ਜ	. P	1675	521			A2													
						B1 20070228				EP 2004-764259									
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		20062										2004 -							
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OTHER	CO	שממו	(C).			MADI	ידי איני	1/2.	2461			-		_		_			

MARPAT 142:246185 OTHER SOURCE(S):

Topical pharmaceutical compns., e.g., an emulsion, comprises a quinazoline derivative (a lavendustin analog), and an emollient, and optionally further excipients. Thus, a formulation contained 1% and iso-Pr myristate 10% in addition to other standard excipients.

182878-97-3 IT

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(pharmaceutical compns. containing quinazoline derivs.)

182878-97-3 CAPLUS RN

Quinazoline, 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl- (CA INDEX NAME)

=> d 17 ibib abs hitstr 2-YOU HAVE REQUESTED DATA FROM 2 ANSWERS - CONTINUE? Y/(N):y

1.7 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2003:662875 CAPLUS

DOCUMENT NUMBER:

139:293651

TITLE:

A Practical Synthesis of 6-[2-(2,5-

Dimethoxyphenyl)ethyl]-4-ethylquinazoline and the Art

of Removing Palladium from the Products of

Pd-Catalyzed Reactions

AUTHOR (S):

Koenigsberger, Kurt; Chen, Guang-Pei; Wu, Raeann R.;

Girgis, Michael J.; Prasad, Kapa; Repic, Oljan;

Blacklock, Thomas J.

CORPORATE SOURCE:

Process Research and Development, Novartis Institute for Biomedical Research, East Hanover, NJ, 07936, USA

SOURCE:

Organic Process Research & Development (2003), 7(5),

733-742

CODEN: OPRDFK; ISSN: 1083-6160

PUBLISHER:

American Chemical Society

DOCUMENT TYPE:

Journal

LANGUAGE:

English

OTHER SOURCE(S):

CASREACT 139:293651

A concise large-scale synthesis of 1, a new antimitotic agent is described. The key step was a one-pot Sonogashira cross-coupling of an aryl halide with a heteroaryl halide through an acetylene using the readily available 2-methyl-3-butyn-2-ol (7). An innovative approach for palladium removal was designed and successfully scaled-up on a multikilogram scale. The product was crystallized from the crude reaction mixture while keeping the residual palladium in the mother liquor by using Pd-scavenging agents such as N-acetylcysteine or thiourea.

182878-97-3P, 6-[2-(2,5-Dimethoxyphenyl)ethyl]-4-ethylquinazoline IT RL: IMF (Industrial manufacture); PUR (Purification or recovery); PREP (Preparation)

> (scale-up of and Pd catalyst removal in the synthesis of 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethylquinazoline)

RN 182878-97-3 CAPLUS

Quinazoline, 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl- (CA INDEX NAME) CN

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

23

ACCESSION NUMBER: DOCUMENT NUMBER:

REFERENCE COUNT:

1996:661128 CAPLUS

125:301017

TITLE:

Preparation of trisubstituted phenyl and quinazoline

THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS

derivatives for the treatment of inflammatory and

proliferative skin diseases and cancer

INVENTOR(S):

Nussbaumer, Peter

PATENT ASSIGNEE(S):

Sandoz Ltd., Switz.; Sandoz-Patent-Gmbh;

Sandoz-Erfindungen Verwaltungsgesellschaft Mbh

SOURCE: PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

						APPLICATION NO.						DATE						
WO									WO 1996-EP1116									
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		LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO	, NZ	, PL	, PT	, RO,	RU,	SD,	SE,	
		SG,	SI															
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			B1 20011114															
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RU	21642	224								RU :	1997-	-117	169		1	9960	314	
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AΤ	20876	63			Т		2001	1115		AT :	1996-	- 908	043		1	9960	314	
CZ	28994				В6		2002	0417		CZ :	1997-	-286	9		1	9960	314	
PT	81508				T		2002	0429		PT :	1996	-908	043		1	9960	314	
ES	21684	463														9960	314	
	1858															9960	314	
	97034															9970	821	
	9704															9970	-	
		-									-							

NO 310356	B1	20010625				
US 5990116	A	19991123	US	1997-913597		19970915
HK 1014440	A1	20030627	HK	1998-112775		19981203
PRIORITY APPLN. INFO.:			GB	1995-5080	Α	19950314
			GB	1995-5858	Α	19950323
			GB	1995-26593	A	19951228
			WO	1996-EP1116	W	19960314
OTHER COIDCE(C).	יד אם פאא	125.301017				

OTHER SOURCE(S): MARPAT 125:30101

GΙ

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB The invention concerns compds. I wherein: R1 and R2 are the same or different and represent OH, alkoxy, acyloxy, alkyl, or acyl, whereby R2 is in the 5- or 6-position, with the proviso that R1 and R2 are not simultaneously OH or acyloxy, and (a) W represents CH2CH2, R3 = C(:X)R6 wherein R6 = H, alkyl, alkoxy, or amino and X = O, hydroxyimino, or alkoxyimino, R4 = NR7R8 wherein R7 and R8 are the same or different and represent H, alkyl, acyl, alkoxycarbonyl, or (b) W = CH2CH2, CH:CH, CH2O, or CH2NR5, whereby the heteroatom adheres to ring B and R5 = H, alkyl, or acyl, R3 and R4 form together with ring B a condensed ring system II or III wherein the single/dotted line symbol represents a single or double bond, R9 = e.g., H, alkylthio, alkyl; Y = N or CR11, R10 = e.g., H, alkyl, acyl; R11 = H, alkoxycarbonyl, cyano, acyl; Z = O or S; and V = NH if the single/dotted line represents a single bond, and N if the single/dotted line represents a double bond (with provisos), and their use in the prevention or treatment of inflammatory and proliferative skin diseases and cancer. Thus, e.g., hydrogenation of 6-[2-(2,5dimethoxyphenyl)ethynyl]-4-ethylquinazoline afforded 6-[2-(2,5dimethoxyphenyl)ethyl]-4-ethylquinazoline (IV); IV and the corresponding 4-MeO and 4-Me compds. exhibited IC50 of about 10 nM for the inhibition of proliferation in the human keratinocyte cell line HaCaT, and between 10 and 200 nM for inhibition of tumor cell proliferation.

IT 182878-97-3P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of trisubstituted Ph and quinazoline derivs. for the treatment of inflammatory and proliferative skin diseases and cancer)

RN 182878-97-3 CAPLUS

CN Quinazoline, 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl- (CA INDEX NAME)

WEST Search History

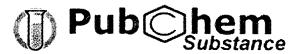
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DATE: Tuesday, November 27, 2007

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Г	L5	((psoriasis and emollient and psoriasis.ti,ab,clm. and emollient.ti,ab,clm.) and (@pd<20030819 or @ad<20030819 or @rlad<20030819 or @prad<20030819))	111
	DB=P	GPB, USPT; THES=ASSIGNEE; PLUR=YES; OP=ADJ	
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	DB=P	GPB, USPT, USOC, EPAB, JPAB, DWPI; THES=ASSIGNEE; PLUR=YES; OP=AD	J
Γ	L3	(psoriasis and emollient)	2377
	DB=P	GPB, USPT; THES=ASSIGNEE; PLUR=YES; OP=ADJ	
	L2	(5990116.pn.)	1
	DB=P	GPB, USPT, USOC, EPAB, JPAB, DWPI; THES=ASSIGNEE; PLUR=YES; OP=AD	Ŋ
Г	L1	("6-[2-(2,5-Dimethoxyphenyl)ethyl]-4-ethylquinazoline" or "6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl-quinazoline" or (182878-97-3))	0

END OF SEARCH HISTORY





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#6 Search 6-[2-(2,5-dimethoxyphenyl)ethyl]-4-ethyl-Quinazoline

#2 Search 6-[2-(2,5-Dimethoxyphenyl)ethyl]-4ethylquinazoline

#1 Search lavendustin

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PubChem Compound Unique structures with

computed properties

Clear History

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